DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration
Rockville MD 20857

FILE COPY

October 29, 1999

Robert S. Milanese
National Association of Pharmaceutical Manufacturers
3279 Veterans Memorial Highway
Suite D-7
Ronkonkoma, NY 11779

Dear Mr. Milanese:

Your petition requesting the Food and Drug Administration to continue to review and approve ANDA suitability petitions seeking a change in strength, dosage form, active ingredient, or route of administration, was received by this office on 10/26/99. It was assigned docket number 99P-4618/CP 1 and it was filed on 10/26/99. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Jennie Butler

Jennie Butler
Dockets Management Branch

99P-4618

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